



**University of  
Zurich**<sup>UZH</sup>

**Zurich Open Repository and  
Archive**

University of Zurich  
University Library  
Strickhofstrasse 39  
CH-8057 Zurich  
[www.zora.uzh.ch](http://www.zora.uzh.ch)

---

Year: 2014

---

## **Assessing the burden of healthcare-associated infections through prevalence studies: what is the best method?**

Zingg, Walter ; Huttner, Benedikt D ; Sax, Hugo ; Pittet, Didier

**Abstract:** **OBJECTIVE:** To explore differences in the prevalence of healthcare-associated infections (HAIs) according to survey methodology. **DESIGN:** Repeated point and period prevalence survey strategies. **SETTING:** University-affiliated primary and tertiary care center. **METHODS:** Analysis of data collected from 2006 to 2012 from annual HAI prevalence surveys using definitions proposed by the US Centers for Disease Control and Prevention. The study design allowed the analysis of the same data in the format of a point or a period prevalence survey. **RESULTS:** Pooled point and period HAI prevalence was 7.46% and 9.84% (+32%), respectively. This additional 32% was mainly attributable to infections of the lower respiratory tract (2.42% vs 3.20% [+32%]) and the urinary tract (1.76% vs 2.62% [+49%]). Differences in surgical site infections (1.02% vs 1.20% [+19%]) and bloodstream infections (0.76% vs 0.86% [+13%]) were smaller. HAI prevalence for the point and period methodology in acute and long-term care were 7.47% versus 9.38 (+26%) and 8.37% versus 11.89% (+42%), respectively. Differences were stable over time. Focusing on the 4 major HAIs (respiratory tract, urinary tract, surgical site, and bloodstream infections) misses one-quarter of all HAIs. **CONCLUSIONS:** More HAIs are identified by the period prevalence method, especially those of shorter duration (lower respiratory and urinary tract), which would make this method more suitable to be used in long-term care. Results of the 2 study methods cannot be benchmarked against each other.

DOI: <https://doi.org/10.1086/676424>

Posted at the Zurich Open Repository and Archive, University of Zurich

ZORA URL: <https://doi.org/10.5167/uzh-102410>

Journal Article

Published Version

Originally published at:

Zingg, Walter; Huttner, Benedikt D; Sax, Hugo; Pittet, Didier (2014). Assessing the burden of healthcare-associated infections through prevalence studies: what is the best method? *Infection Control and Hospital Epidemiology*, 35(6):674-684.

DOI: <https://doi.org/10.1086/676424>



CHICAGO JOURNALS



---

Assessing the Burden of Healthcare-Associated Infections through Prevalence Studies: What Is the Best Method?

Author(s): Walter Zingg, MD; Benedikt D. Huttner, MD, MS; Hugo Sax, MD; Didier Pittet, MD, MS

Source: *Infection Control and Hospital Epidemiology*, Vol. 35, No. 6 (June 2014), pp. 674-684

Published by: [The University of Chicago Press](#) on behalf of [The Society for Healthcare Epidemiology of America](#)

Stable URL: <http://www.jstor.org/stable/10.1086/676424>

Accessed: 16/12/2014 11:01

---

Your use of the JSTOR archive indicates your acceptance of the Terms & Conditions of Use, available at <http://www.jstor.org/page/info/about/policies/terms.jsp>

JSTOR is a not-for-profit service that helps scholars, researchers, and students discover, use, and build upon a wide range of content in a trusted digital archive. We use information technology and tools to increase productivity and facilitate new forms of scholarship. For more information about JSTOR, please contact support@jstor.org.



*The University of Chicago Press* and *The Society for Healthcare Epidemiology of America* are collaborating with JSTOR to digitize, preserve and extend access to *Infection Control and Hospital Epidemiology*.

<http://www.jstor.org>

## ORIGINAL ARTICLE

# Assessing the Burden of Healthcare-Associated Infections through Prevalence Studies: What Is the Best Method?

Walter Zingg, MD;<sup>1</sup> Benedikt D. Huttner, MD, MS;<sup>1</sup> Hugo Sax, MD;<sup>1,a</sup> Didier Pittet, MD, MS<sup>1</sup>

**OBJECTIVE.** To explore differences in the prevalence of healthcare-associated infections (HAIs) according to survey methodology.

**DESIGN.** Repeated point and period prevalence survey strategies.

**SETTING.** University-affiliated primary and tertiary care center.

**METHODS.** Analysis of data collected from 2006 to 2012 from annual HAI prevalence surveys using definitions proposed by the US Centers for Disease Control and Prevention. The study design allowed the analysis of the same data in the format of a point or a period prevalence survey.

**RESULTS.** Pooled point and period HAI prevalence was 7.46% and 9.84% (+32%), respectively. This additional 32% was mainly attributable to infections of the lower respiratory tract (2.42% vs 3.20% [+32%]) and the urinary tract (1.76% vs 2.62% [+49%]). Differences in surgical site infections (1.02% vs 1.20% [+19%]) and bloodstream infections (0.76% vs 0.86% [+13%]) were smaller. HAI prevalence for the point and period methodology in acute and long-term care were 7.47% versus 9.38 (+26%) and 8.37% versus 11.89% (+42%), respectively. Differences were stable over time. Focusing on the 4 major HAIs (respiratory tract, urinary tract, surgical site, and bloodstream infections) misses one-quarter of all HAIs.

**CONCLUSIONS.** More HAIs are identified by the period prevalence method, especially those of shorter duration (lower respiratory and urinary tract), which would make this method more suitable to be used in long-term care. Results of the 2 study methods cannot be benchmarked against each other.

*Infect Control Hosp Epidemiol* 2014;35(6):674-684

The pioneering Study on the Efficacy of Nosocomial Infection Control (SENIC) project, initiated in the 1970s by the US Centers for Disease Control and Prevention (CDC), unequivocally proved the benefit of healthcare-associated infection (HAI) surveillance.<sup>1,2</sup> The method was based on a stratified random sample of patients from 338 US hospitals, and HAIs were detected by thorough patient chart review. The HAI prevalence at that time was estimated at approximately 5.2%.<sup>3-5</sup> In 1970, the US National Nosocomial Infection Surveillance network was established to provide regular prospective outcome data on HAI in intensive care units (ICUs) in the United States.<sup>6</sup> In parallel, the CDC issued definitions of nosocomial infections.<sup>7</sup> Over the following decades, the CDC HAI definitions were continually updated and became the reference standard for the vast majority of HAI surveillance activities around the world.<sup>7-14</sup>

In 1981, the World Health Organization (WHO) convened an advisory group on the surveillance, control, and prevention of HAI.<sup>15</sup> The group specifically recommended the con-

duct of HAI prevalence surveys to assess the burden of the problem in different parts of the world. Later, WHO published prevalence data gathered between 1983 and 1985 from 47 hospitals in 14 countries.<sup>15</sup> At the same time, an increasing number of countries started to conduct national or regional prevalence surveys. Most recently, the European Centre for Disease Prevention and Control (ECDC), as well as the CDC, performed large point prevalence surveys in Europe and the United States based on the methodology published in 2 pilot studies, and the results of the ECDC point prevalence survey are now published.<sup>16-18</sup> Most local, regional, and national surveys used the point prevalence methodology, that is, only HAIs active on the day of the survey were taken into account. However, some studies in Italy,<sup>19,20</sup> Switzerland,<sup>21-25</sup> and the United States<sup>26</sup> used the period prevalence method, that is, not only were HAIs active on the day of the survey accounted for, but those active during a predefined period before the survey day were also assessed (Figure 1). Some surveys, such as the first Spanish prevalence survey of the Estudio de Prev-

**Affiliations:** 1. Infection Control Program and World Health Organization Collaborating Center on Patient Safety, University of Geneva Hospitals and Faculty of Medicine, Geneva, Switzerland; a. Present affiliation: Division of Infectious Diseases and Infection Control, University Hospital of Zurich and Faculty of Medicine, Zurich, Switzerland.

Received November 1, 2013; accepted January 8, 2014; electronically published April 17, 2014.

© 2014 by The Society for Healthcare Epidemiology of America. All rights reserved. 0899-823X/2014/3506-0007\$15.00. DOI: 10.1086/676424

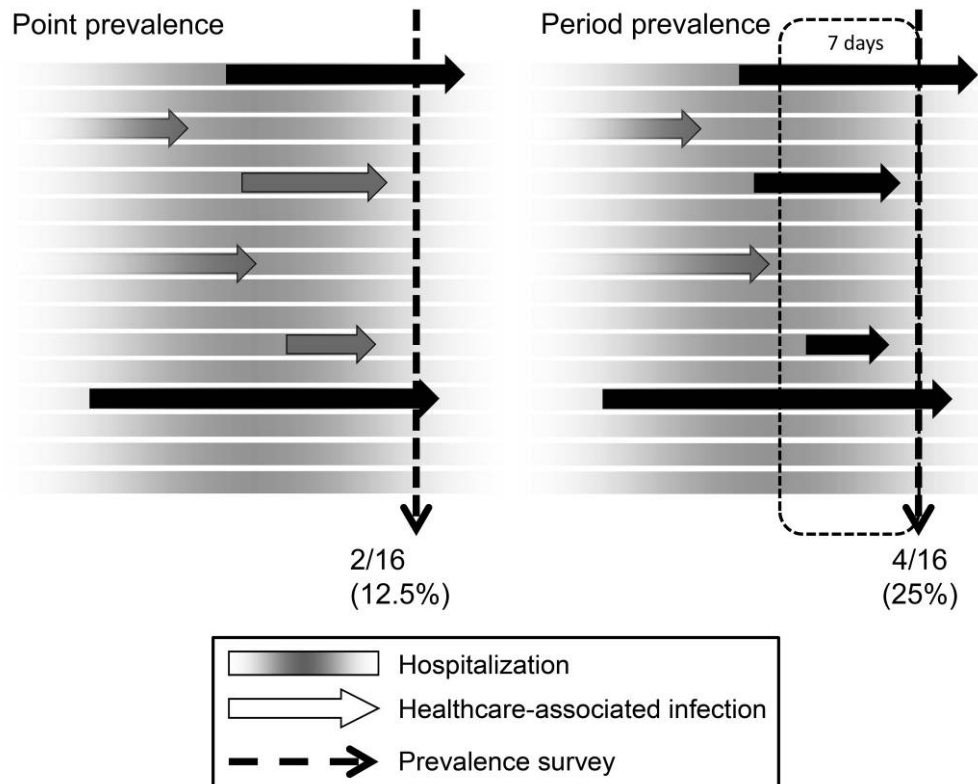


FIGURE 1. The concepts of point and period prevalence surveys. In point prevalence surveys, all patients present on the day of prevalence are eligible, and only healthcare-associated infections active at the time of the survey are included in the analysis. In this example, 2 of 16 patients have a healthcare-associated infection, for a prevalence of 12.5%. In period prevalence surveys, all patients present on the day of prevalence are eligible, and healthcare-associated infections active at the time of the survey or in the preceding 6 days (for a total of 7 days of study) are included in the analysis. In this example, 4 of 16 patients have a healthcare-associated infection, for a prevalence of 25%.

alencia de las Infecciones Nosocomiales en Espana (EPINE) network, combined point prevalence (active infection on the day of survey) with extrinsic risk factors present in the 7 days before the survey.<sup>27</sup> Both methodologies have advantages and disadvantages. While a period prevalence survey will allow capture of more HAIs, especially those of short duration, it is methodologically “less pure,” since it mixes the concepts of prevalence and incidence, and it is also more time-consuming than a pure point prevalence survey.

To our knowledge, no study has analyzed the differences between the point and period methodology to assess the burden of HAI. We conducted this study to provide such an analysis from a large database and to place the findings in the context of the published literature.

## METHODS

The University of Geneva Hospitals (Geneva, Switzerland) is a primary and tertiary care center with 1,908 beds; in 2012, 47,000 admitted patients accumulated 670,000 patient-days. Located at 8 different sites, the hospital offers intensive, acute, and long-term inpatient care and also includes a pediatric hospital. Since 1994, the infection control team has conducted

annual period prevalence surveys in May and early June.<sup>28</sup> The current study focused on data collected from 2006 to 2012 from all departments except psychiatry. All types of HAI as defined by the CDC were included apart from asymptomatic urinary tract infection (UTI).<sup>11,12</sup> No adjustments or modifications of the definitions or other aspects of the methodology were made during the study period, and all staff involved in the survey were trained in the methodology.

Every patient present in the ward on the day of the prevalence survey was included except those admitted on the calendar day of the survey. Infection control nurses and physicians screened patient charts for clinical symptoms and signs, laboratory data, microbiological results, and information from other diagnostics suggestive for infection within a 1-week period, ending with the day of the prevalence survey. HAIs were counted when they were active at any time within the 1-week period (Figure 1).<sup>28</sup> An infection was considered active when the patient had clinical symptoms and/or was still receiving treatment for that infection. Surgical site infections were documented as healthcare associated when they occurred within 30 days after the operation or 1 year in the case of infection associated with the insertion of a prosthetic

TABLE 1. Patient Characteristics with Annual Trends: Prevalence Surveys, University of Geneva Hospitals, 2006–2012

| Characteristic                            | Pooled data     | Trend, IRR (95% CI) |
|---|-----------------|---------------------|
| Age, median (IQR), years                  | 72 (50–83)      | 1.01 (1.01–1.01)    |
| Sex (female)                              | 5,774 (56)      | 1.00 (0.99–1.02)    |
| Charlson comorbidity index, mean $\pm$ SD | 1.17 $\pm$ 1.68 | 1.01 (1.00–1.02)    |
| McCabe score, mean $\pm$ SD               | 1.19 $\pm$ 0.46 | 1.01 (1.00–1.02)    |
| Surgery                                   | 2,554 (25)      | 1.01 (0.99–1.03)    |
| ICU stay at any time                      | 853 (8)         | 1.07 (1.03–1.10)    |
| Distribution of acute and long-term care  |                 |                     |
| Acute care at prevalence                  | 5,717 (55)      | 1.00 (0.99–1.02)    |
| Long-term care at prevalence              | 4,650 (45)      | 1.00 (0.98–1.01)    |

NOTE. Data are no. (%), unless otherwise indicated. CI, confidence interval; ICU, intensive care unit; IQR, interquartile range; IRR, incidence rate ratio; SD, standard deviation.

device.<sup>21</sup> The distinction between point and period prevalence was possible because the data set contained a variable indicating whether an HAI was active on the day of the prevalence survey. Annual prevalence surveys are part of a quality improvement program promoted by the directorate of the University of Geneva Hospitals. The institutional ethics committee waived informed consent related to the prevalence surveys. HAI duration was estimated from the pooled data by the difference of days between the date of HAI onset and the date of the point prevalence survey.

PubMed was searched for published prevalence surveys without restrictions of language up to June 30, 2013. The following search term was used: (“prevalence” [Title] OR “point-prevalence” [Title] OR “cross-sectional” [Title]) AND (“nosocomial” [Title] OR “hospital-acquired” [Title] OR “infection in hospitals” [Title] OR “hospital infection” [Title] OR “hospital infections” [Title] OR “healthcare-associated” [Title] OR “hospital-associated” [Title] OR “infections among hospitalized patients” [Title]). Further references were obtained by a full text sift of retrieved publications.

### Statistical Analysis

The descriptive analysis was stratified by CDC infection categories (lower respiratory tract infection and pneumonia [LRTI], UTI, surgical site infection [SSI], bloodstream infection and clinical sepsis [BSI], gastrointestinal infection [GI], skin and soft-tissue infection [SST], and other infections, which included mostly eye, ear, nose, and throat infections) and care settings (acute [including intensive care] and long-term care). Descriptive statistics were used to express the difference between point and period prevalence surveys. No formal statistical test was used to quantify the differences because the same database was used to calculate outcomes for both point and period prevalence (the null hypothesis that the prevalence obtained by the different methodologies is equivalent can be rejected without a test since the period prevalence contains all infections counted in the point prevalence). Differences for the different HAIs between acute and long-term care were analyzed using a simple  $\chi^2$  test. Trends

related to patient characteristics across the study years were determined using a nonadjusted Poisson regression analysis for each separate variable and reported as incidence rate ratios. Two-sided  $\alpha < .05$  was used to determine statistical significance. All statistical analyses were conducted using Stata software, version 10.0 (StataCorp).

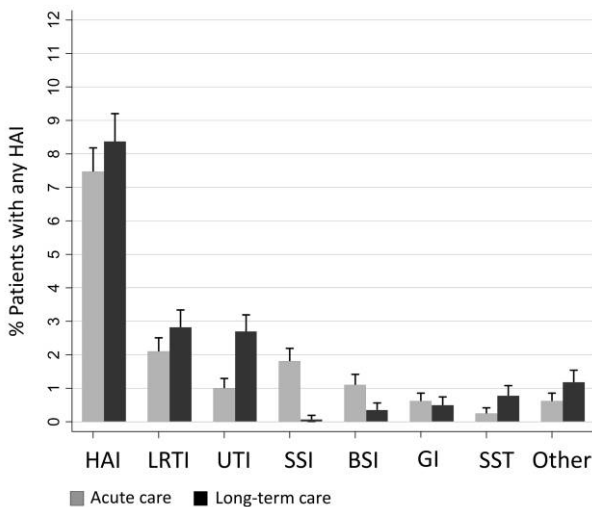
### RESULTS

A total of 7 annual prevalence surveys including 10,367 patients were analyzed. Patient characteristics such as sex, age, Charlson comorbidity index, surgery, and distribution among the different care settings are summarized in Table 1. We observed moderate yearly trends toward higher age, higher McCabe classification<sup>29</sup> and Charlson comorbidity index<sup>30</sup> scores, and more frequent ICU stays. Proportions of acute and long-term care did not change during the study period (Table 1).

The point and period prevalence surveys identified a total of 816 and 1,089 HAIs among 773 and 1,020 patients, respectively. Estimated HAI durations (median [interquartile range]) for LRTI, UTI, SSI, BSI, GI, SST, and other infections were 6 (3–10) days, 5 (3–9) days, 14.5 (6–29) days, 7 (3–12) days, 6.5 (3–11) days, 6 (4–13) days, and 7.5 (5–15) days, respectively.

The pooled point and period prevalence (95% confidence interval) of all HAIs were 7.46% (6.96%–7.98%) and 9.84% (9.27%–10.42%), respectively. Figure 2 summarizes the differences of pooled HAI prevalence between acute and long-term care. Significantly higher proportions of UTI and LRTI were identified in long-term care, while the proportions of SSI and BSI were higher in acute care, irrespective of the prevalence methodology. Table 2 summarizes the differences between pooled point and period prevalence surveys for HAI, LRTI, UTI, SSI, BSI, GI, SST, and other infections stratified into acute and long-term care. Overall, HAI ratios were higher (+33.4%) when assessed by the period prevalence method in both acute (+25.6%) and long-term (+42.1%) care. There were particularly large differences between the 2 methodol-

### A. Point prevalence



### B. Period prevalence

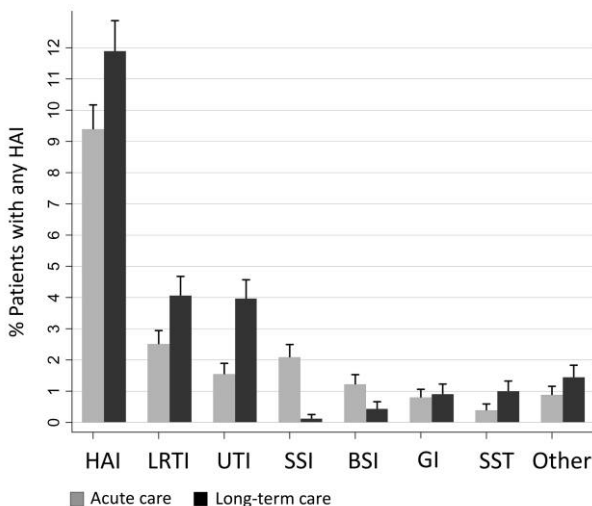


FIGURE 2. Pooled point and period prevalence of healthcare-associated infections (HAIs) with 95% confidence intervals stratified by clinical setting: prevalence surveys, University of Geneva Hospitals, 2006–2012. A, Point prevalence. B, Period prevalence. BSI, bloodstream infection; GI, gastrointestinal infection; LRTI, pneumonia/lower respiratory tract infection; SSI, surgical site infection; SST, skin and soft-tissue infection; UTI, urinary tract infection.

ogies for LRTI and UTI in long-term care where more of these infections were identified. SSIs and BSIs were identified mostly in acute care with little difference between point and period prevalence.

Focusing on the 4 leading HAIs (LRTI, UTI, SSI, and BSI) misses a high proportion of infections in both the point (24.3%) and the period prevalence (25.0%), respectively. Differences are larger in long-term than acute care in both the point (33.0% vs 19.7%) and the period prevalence (28.0% vs

21.9%). The difference between the 2 settings is more pronounced in the point than in the period prevalence.

### Results of the Literature Search

The search term identified 305 publications, of which 249 were prevalence surveys or complementary material. An additional 4 studies were identified by a search of the references of retrieved publications. A total of 97 international, national, or regional multicenter (more than 1 center) published surveys were identified with an upward trend over the decades.

### DISCUSSION

Our study shows that benchmarking between point and period prevalence data is not possible, as the higher proportion of infections identified by the period methodology favors HAIs of short duration and infections in long-term care. Prevalence surveys are biased in favor of HAIs of longer duration compared with incidence surveys and are notably influenced by the duration of antimicrobial treatment and the propensity to discharge patients.<sup>4</sup> The period prevalence methodology counterbalances this to some degree, and the proportion of the different HAIs becomes more similar to the proportion of their incidence.

From a methodological point of view, the idea of the period prevalence may be challenged because it mixes the concepts of prevalence and incidence (J. Freeman, personal communication, 1999). However, the period methodology should not be considered inferior to the point prevalence methodology on the basis of this aspect alone, and our data suggest that it could be of particular interest for use in long-term care settings where HAIs of short duration, such as LRTI and UTI, are common and where the burden of these HAIs may be underestimated when using the point prevalence method. The difference between the 2 methodologies in acute care settings, where more SSIs and BSIs occur, is less important. Period prevalence data do not serve to estimate incidence rates using common algorithms.<sup>31–33</sup> The better approximation of HAIs of short duration by the period methodology could add value to such models, but further studies are needed to validate this hypothesis. Furthermore, focusing on the 4 leading (ie, most frequent) HAIs (LRTI, UTI, SSI, and BSI) may reduce workload but underestimates the proportion of infections (such as GI, SST, eye, and ear, nose, and throat infections) that are common in long-term care. This makes such an approach unsuitable for this type of setting, and prevalence surveys in long-term care should better focus on a more appropriate selection of HAIs, such as LRTI, UTI, SSI, and SST<sup>34</sup> or LRTI, UTI, BSI, GI, and conjunctivitis.<sup>35</sup>

HAI incidence data have become the gold standard of HAI surveillance over the past 2 decades. National and multinational networks, such as the US National Healthcare Safety Network, the German Krankenhaus Infektions Surveillance System, and the International Nosocomial Infection Control Consortium, have become success stories both in high-income



TABLE 2. Distribution of Pooled Point and Period Prevalence of Healthcare-Associated Infections (HAIs) Stratified by Clinical Setting: Prevalence Surveys, University of Geneva Hospitals, 2006–2012

| Clinical setting, type of infection | Point prevalence, <sup>a</sup> |                  | Period prevalence, <sup>a</sup> |                     |
|-------------------------------------|--------------------------------|------------------|---------------------------------|---------------------|
|                                     | No. (%)                        | % (95% CI)       | No. (%)                         | % (95% CI)          |
| Hospital-wide (10,367 patients)     |                                |                  |                                 |                     |
| LRTI                                | 251 (31)                       | 2.42 (2.13–2.74) | 332 (30)                        | 3.20 (2.87–3.56)    |
| UTI                                 | 182 (22)                       | 1.76 (1.51–2.03) | 272 (25)                        | 2.62 (2.32–2.95)    |
| SSI                                 | 106 (13)                       | 1.02 (0.84–1.24) | 124 (11)                        | 1.20 (1.00–1.42)    |
| BSI                                 | 79 (10)                        | 0.76 (0.60–0.95) | 89 (8)                          | 0.86 (0.69–1.06)    |
| GI                                  | 58 (7)                         | 0.56 (0.43–0.72) | 87 (8)                          | 0.84 (0.67–1.03)    |
| SST                                 | 50 (6)                         | 0.48 (0.36–0.64) | 68 (6)                          | 0.66 (0.51–0.83)    |
| Other                               | 90 (11)                        | 0.87 (0.70–1.07) | 117 (11)                        | 1.13 (0.93–1.35)    |
| All HAI                             | 816 (100)                      | 7.87 (7.36–8.41) | 1,089 (100)                     | 10.50 (9.92–11.11)  |
| Acute care (5,717 patients)         |                                |                  |                                 |                     |
| LRTI                                | 120 (28)                       | 2.10 (1.74–2.50) | 143 (27)                        | 2.50 (2.11–2.94)    |
| UTI                                 | 57 (13)                        | 1.00 (0.76–1.29) | 88 (16)                         | 1.54 (1.24–1.89)    |
| SSI                                 | 103 (24)                       | 1.80 (1.47–2.19) | 119 (22)                        | 2.08 (1.73–2.49)    |
| BSI                                 | 63 (15)                        | 1.10 (0.85–1.41) | 69 (13)                         | 1.21 (0.94–1.52)    |
| GI                                  | 35 (8)                         | 0.61 (0.43–0.85) | 45 (8)                          | 0.79 (0.57–1.05)    |
| SST                                 | 14 (3)                         | 0.24 (0.13–0.41) | 22 (4)                          | 0.38 (0.24–0.58)    |
| Other                               | 35 (8)                         | 0.61 (0.43–0.85) | 50 (9)                          | 0.87 (0.65–1.15)    |
| All HAI                             | 427 (100)                      | 7.47 (6.80–8.18) | 536 (100)                       | 9.38 (8.63–10.16)   |
| Long-term care (4,650 patients)     |                                |                  |                                 |                     |
| LRTI                                | 131 (34)                       | 2.82 (2.36–3.33) | 189 (34)                        | 4.06 (3.52–4.67)    |
| UTI                                 | 125 (32)                       | 2.69 (2.24–3.19) | 184 (33)                        | 3.96 (3.42–4.56)    |
| SSI                                 | 3 (1)                          | 0.06 (0.01–0.19) | 5 (1)                           | 0.11 (0.03–0.25)    |
| BSI                                 | 16 (4)                         | 0.34 (0.20–0.56) | 20 (4)                          | 0.43 (0.26–0.66)    |
| GI                                  | 23 (6)                         | 0.49 (0.31–0.74) | 42 (8)                          | 0.90 (0.65–1.22)    |
| SST                                 | 36 (9)                         | 0.77 (0.54–1.07) | 46 (8)                          | 0.99 (0.72–1.32)    |
| Other                               | 55 (14)                        | 1.18 (0.89–1.54) | 67 (12)                         | 1.44 (1.12–1.83)    |
| All HAI                             | 389 (100)                      | 8.37 (7.59–9.20) | 553 (100)                       | 11.89 (10.98–12.86) |

NOTE. CI, confidence interval; BSI, bloodstream infection; GI, gastrointestinal infection; LRTI, pneumonia/lower respiratory tract infection; SSI, surgical site infection; SST, skin and soft-tissue infection; UTI, urinary tract infection.

<sup>a</sup> Infections divided by the total number of patients.

and in low- and middle-income countries, thus serving as a reference for many similar undertakings.<sup>36–38</sup> Prospective hospital-wide HAI incidence surveillance is time-consuming and costly, and such programs are often restricted to the ICU or other high-risk settings, such as oncology or neonatology. However, HAI is not exclusively confined to such high-risk areas; it also occurs in regular wards. For example, it has been shown that central line-associated BSIs occur at similar incidence rate ratios both in and outside the ICU.<sup>39</sup>

Prevalence surveys help to assess the burden of hospital-wide HAI at a reasonable cost.<sup>40–42</sup> In the 1970s, many countries and regions in Europe and elsewhere began to conduct prevalence surveys, and the CDC and the ECDC have now followed this example by launching large point prevalence surveys in the United States and Europe.<sup>18</sup> Table 3 and Figure 3 show when countries and regions started to conduct international, national, or regional prevalence surveys, spanning a timeline of 4 decades. Only a few prevalence surveys used the period methodology,<sup>19–26,43</sup> and the vast majority in Africa,<sup>44–50</sup> Asia,<sup>51–64</sup> Australasia,<sup>65–67</sup> Europe,<sup>34,40,42,68–110</sup> North

America,<sup>17,111–113</sup> and South America<sup>114–117</sup> used the point prevalence methodology. Although many surveys used some version of the CDC HAI definitions, methodological inequalities make direct comparison of prevalence results difficult.<sup>118</sup>

The history of HAI prevalence surveys is linked to the meticulous work of the SENIC project in the 1970s and the useful HAI definitions issued and regularly updated by the CDC and the ECDC, which have served the infection control community for many years. Today, time and budget restrictions force infection control programs to reestablish the concept of the prevalence survey, despite its methodological limitations. The recent commitment of the CDC and the ECDC to prevalence surveys sets the stage for future prevalence studies using comparable methodologies. More than 30 years after it was issued, the WHO call for action to perform HAI prevalence surveys to assess the size of the problem in different parts of the world is finally established.<sup>15</sup>

In summary, our findings suggest that the additional portion of HAIs detected by the period methodology favors HAIs of short duration, which are less likely to be captured by the

TABLE 3. First International, National, or Regional Prevalence Surveys of Healthcare-Associated Infections: 1970–2010

| Country/organization          | Year(s)<br>of survey | Hospitals       | Patients | Setting(s) | CDC criteria           | Method             | Prevalence, <sup>a</sup> % |
|-------------------------------|----------------------|-----------------|----------|------------|------------------------|--------------------|----------------------------|
| Sweden <sup>69</sup>          | 1975                 | 5               | 4,246    | All        | Yes <sup>7</sup>       | Point              | 17.0                       |
| Denmark <sup>70</sup>         | 1978                 | 20/25           | 2,920    | Acute      | Yes <sup>8</sup>       | Point              | 10.4                       |
| United Kingdom <sup>75</sup>  | 1979                 | 43              | 18,163   | Acute      | Yes <sup>10</sup>      | Point              | 19.1                       |
| Italy <sup>68</sup>           | 1983                 | 130             | 34,577   | Acute      | Yes <sup>8</sup>       | Point              | 19.3                       |
| WHO <sup>15</sup>             | 1983–1985            | 47              | 28,861   | Acute      | No                     | Point              | 8.7                        |
| Australia <sup>65</sup>       | 1984                 | 269             | 28,643   | Acute      | Yes <sup>10,119</sup>  | Point              | 6.3                        |
| Belgium <sup>76</sup>         | 1984                 | 106             | 8,723    | Acute      | Yes <sup>8</sup>       | Point <sup>b</sup> | 9.3                        |
| Czechoslovakia <sup>105</sup> | 1984                 | 23              | 12,260   | All        | No <sup>120</sup>      | Point              | 6.1                        |
| Thailand <sup>53</sup>        | 1988                 | 23              | 6,805    | Acute      | No <sup>121</sup>      | Point              | 11.7                       |
| Spain <sup>27,122</sup>       | 1990                 | 123             | 38,489   | Acute      | Yes <sup>11</sup>      | Point <sup>c</sup> | 8.5                        |
| Norway <sup>78</sup>          | 1991                 | 76              | 14,977   | Acute      | Yes <sup>9</sup>       | Point <sup>d</sup> | 6.3                        |
| Brazil <sup>114</sup>         | 1992                 | 11              | 2,339    | Acute      | Yes <sup>11</sup>      | Point              | 14.0                       |
| Europe <sup>85</sup>          | 1992                 | 1,417           | 10,038   | ICU        | Yes <sup>11</sup>      | Point              | 44.8                       |
| Mauritius <sup>50</sup>       | 1992                 | 4               | 1,190    | Acute      | Yes <sup>10</sup>      | Point              | 4.9                        |
| Germany <sup>94</sup>         | 1995                 | 72              | 14,996   | Acute      | Yes <sup>11</sup>      | Point              | 3.5                        |
| France <sup>82</sup>          | 1996 <sup>e</sup>    | 830             | 236,334  | All        | Yes <sup>14</sup>      | Point              | 6.7                        |
| New Zealand <sup>66,67</sup>  | 1996–1999            | 4               | 5,819    | All        | Yes <sup>11</sup>      | Point              | 9.5                        |
| Switzerland <sup>21</sup>     | 1996                 | 4               | 1,349    | Acute      | Yes <sup>11</sup>      | Period             | 11.6                       |
| Cuba <sup>116</sup>           | 1997                 | 28              | 6,152    | Acute      | Yes <sup>11,14</sup>   | Point              | 6.8                        |
| Lebanon <sup>62</sup>         | 1997                 | 14              | 834      | Acute      | Yes <sup>119,123</sup> | Point              | 6.8                        |
| Greece <sup>102</sup>         | 1999                 | 14              | 3,925    | Acute      | Yes <sup>11,14</sup>   | Point              | 9.3                        |
| Mexico <sup>112</sup>         | 1999 <sup>f</sup>    | 21              | 1,183    | Pediatric  | Yes                    | Point              | 9.8                        |
| Slovenia <sup>106</sup>       | 2001                 | 19              | 6,695    | Acute      | Yes <sup>11,14</sup>   | Point              | 4.6                        |
| Turkey <sup>103</sup>         | 2001                 | 22              | 236      | ICU        | Yes <sup>11</sup>      | Point              | 48.7                       |
| Canada <sup>113</sup>         | 2002                 | 25              | 5,750    | Acute      | Yes                    | Point <sup>g</sup> | 10.5                       |
| Indonesia <sup>63</sup>       | 2001–2002            | 2               | 888      | Acute      | Yes <sup>11,14</sup>   | Point <sup>h</sup> | 8.3                        |
| Latvia <sup>98</sup>          | 2003 <sup>f</sup>    | 2               | 1,291    | Acute      | NA                     | Point              | 5.6                        |
| Iran <sup>60</sup>            | 2004–2005            | 8               | 2,667    | All        | Yes <sup>11</sup>      | Point              | 8.8                        |
| Finland <sup>80</sup>         | 2005                 | 30              | 8,234    | Acute      | Yes <sup>11</sup>      | Point              | 8.5                        |
| Scotland <sup>491</sup>       | 2005–2006            | 45 <sup>i</sup> | 11,608   | Acute      | Yes                    | Point              | 9.5                        |
| Ireland <sup>87</sup>         | 2006                 | 44              | 7,541    | Acute      | Yes <sup>13</sup>      | Point              | 4.9                        |
| China <sup>61</sup>           | 2007–2008            | 13              | 20,350   | Acute      | Yes <sup>11,j</sup>    | Point              | 3.9                        |
| Netherlands <sup>96</sup>     | 2007–2008            | 41              | 26,937   | Acute      | Yes <sup>k</sup>       | Point <sup>l</sup> | 6.2                        |
| Argentina <sup>117</sup>      | 2008                 | 39              | 4,249    | Acute      | Yes <sup>124</sup>     | Point              | 11.3                       |
| Mongolia <sup>58</sup>        | 2008                 | 2               | 933      | Acute      | Yes                    | Point              | 5.4                        |
| Vietnam <sup>56</sup>         | 2008                 | 36              | 7,571    | Acute      | Yes <sup>11</sup>      | Point              | 7.8                        |
| CDC <sup>17</sup>             | 2009                 | 9               | 851      | Acute      | Yes                    | Point              | 6.0                        |
| ECDC <sup>18</sup>            | 2013                 | 947             | 273,753  | Acute      | Yes <sup>12,m</sup>    | Point              | 6.0                        |

NOTE. BSI, bloodstream infection; CDC, US Centers for Disease Control and Prevention; CDI, *Clostridium difficile* infection; ECDC, European Centre for Disease Prevention and Control; ICU, intensive care unit; NA, not available; SSI, surgical site infection; UTI, urinary tract infection; WHO, World Health Organization.

<sup>a</sup> Proportion of patients with 1 or more healthcare-associated infections as defined in the survey.

<sup>b</sup> Only UTI, SSI, and BSI.

<sup>c</sup> Extrinsic risk factors were screened for 7 days before the survey.

<sup>d</sup> All diagnoses except UTIs were based on clinical criteria alone.

<sup>e</sup> A first study among 11,599 patients in 39 hospitals was performed in 1990.<sup>125</sup>

<sup>f</sup> Year of publication.

<sup>g</sup> Only UTI, BSI, SSI, and CDI.

<sup>h</sup> Only UTI, BSI, SSI, and phlebitis.

<sup>i</sup> Acute care only.

<sup>j</sup> With some modifications for infants.

<sup>k</sup> With some modifications.

<sup>l</sup> Only UTI, pneumonia, BSI, and SSI.

<sup>m</sup> European case definitions by HELICS or other European projects were used where available.



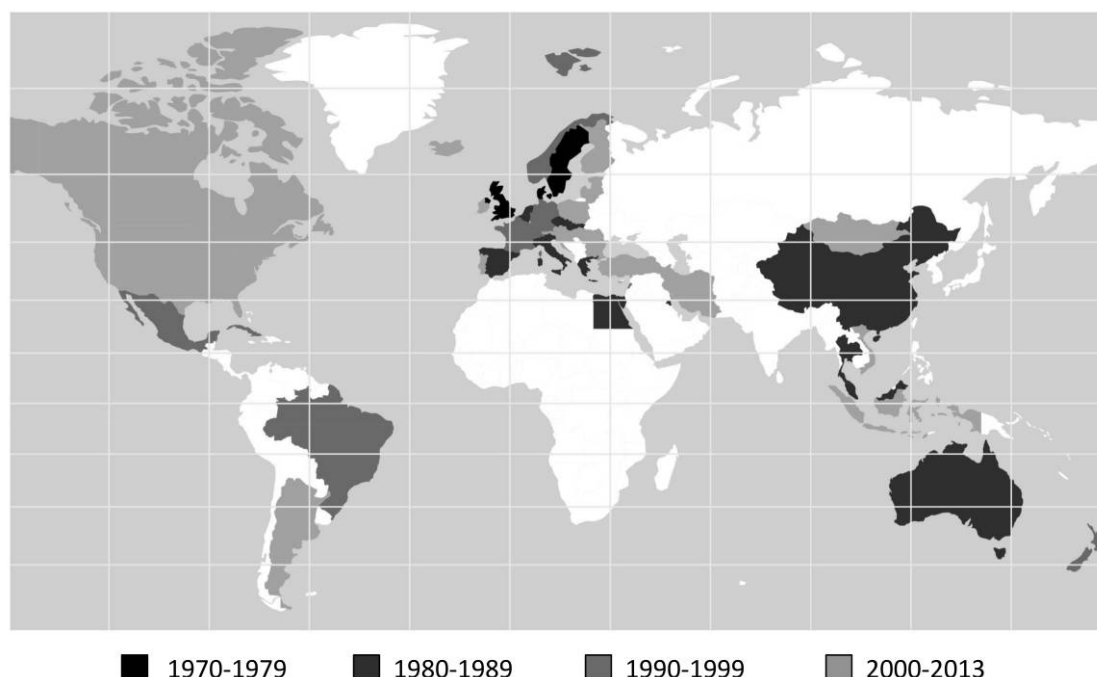


FIGURE 3. International, national, or regional prevalence surveys of healthcare-associated infections in acute or mixed care settings: 1970–2013.

point methodology. This would make the period strategy suitable to be used in long-term care, while the benefit in acute care is low. The results of the 2 concepts cannot be benchmarked against each other. Given the potential advantage of the period methodology in long-term care and the heterogeneity of care settings in our primary and tertiary care hospital, we will continue to use the period methodology with the opportunity to calculate point prevalence data in order to benchmark with other databases.

## ACKNOWLEDGMENTS

We thank the infection control team of the University of Geneva Hospitals for conduct of the surveys, Rosemary Sudan for editorial assistance, and Josep Vaqué and Jonathan Freeman for personal advice.

**Financial support.** The project was part of a quality improvement program financed by the University of Geneva Hospitals.

**Potential conflicts of interest.** All authors report no conflicts of interest relevant to this article. All authors submitted the ICMJE Form for Disclosure of Potential Conflicts of Interest, and the conflicts that the editors consider relevant to this article are disclosed here.

Address correspondence to Walter Zingg, MD, Infection Control Program, University of Geneva Hospitals, 4 Rue Gabrielle Perret-Gentil, 1211 Geneva 14, Switzerland (walter.zingg@hcuge.ch).

Presented in part: 2nd International Conference on Prevention and Infection Control; Geneva, Switzerland; June 25–28, 2013 (Abstract P-222).

## REFERENCES

1. Haley RW, Culver DH, Morgan WM, White JW, Emori TG, Hooton TM. Increased recognition of infectious diseases in US hospitals through increased use of diagnostic tests, 1970–1976. *Am J Epidemiol* 1985;121:168–181.
2. Haley RW, Culver DH, White JW, et al. The efficacy of infection surveillance and control programs in preventing nosocomial infections in US hospitals. *Am J Epidemiol* 1985;121:182–205.
3. Weinstein RA. Nosocomial infection update. *Emerg Infect Dis* 1998;4:416–420.
4. Llata E, Gaynes RP, Fridkin S. Measuring the scope and magnitude of hospital-associated infection in the United States: the value of prevalence surveys. *Clin Infect Dis* 2009;48:1434–1440.
5. Haley RW, Hooton TM, Culver DH, et al. Nosocomial infections in U.S. hospitals, 1975–1976: estimated frequency by selected characteristics of patients. *Am J Med* 1981;70:947–959.
6. Horan TC, White JW, Jarvis WR, et al. Nosocomial infection surveillance, 1984. *MMWR CDC Surveill Summ* 1986;35:17SS–29SS.
7. Centers for Disease Control. *Outline for Surveillance and Control of Nosocomial Infections*. Atlanta: Public Health Service, 1970.
8. Centers for Disease Control. *Outline for Surveillance and Control of Nosocomial Infections*. Atlanta: Public Health Service, 1972.
9. Centers for Disease Control. *Outline for Surveillance and Control of Nosocomial Infections*. Atlanta: Public Health Service, 1976.
10. Garner JS, Bennett JV, Scheckler WE, Maki DG, Brachman PS. Surveillance of nosocomial infections. In: Brachman PS, Eickhoff TC, eds. *International Conference on Nosocomial Infections*. Chicago, 1971:277–281.
11. Garner JS, Jarvis WR, Emori TG, Horan TC, Hughes JM. CDC

- definitions for nosocomial infections, 1988. *Am J Infect Control* 1988;16:128–140.
12. Horan TC, Andrus M, Dudeck MA. CDC/NHSN surveillance definition of health care-associated infection and criteria for specific types of infections in the acute care setting. *Am J Infect Control* 2008;36:309–332.
  13. Horan TC, Gaynes RP. Surveillance of nosocomial infections. In: Mayhall CG, ed. *Hospital Epidemiology and Infection Control*. 3rd ed. Philadelphia: Lippincott Williams & Wilkins, 2004: 1659–1702.
  14. Horan TC, Gaynes RP, Martone WJ, Jarvis WR, Emori TG. CDC definitions of nosocomial surgical site infections, 1992: a modification of CDC definitions of surgical wound infections. *Am J Infect Control* 1992;20:271–274.
  15. Mayon-White RT, Duce G, Kereselidze T, Tikomirov E. An international survey of the prevalence of hospital-acquired infection. *J Hosp Infect* 1988;11(suppl A):43–48.
  16. Zarb P, Coignard B, Griskeviciene J, et al. The European Centre for Disease Prevention and Control (ECDC) pilot point prevalence survey of healthcare-associated infections and antimicrobial use. *Euro Surveill* 2012;17.
  17. Magill SS, Hellinger W, Cohen J, et al. Prevalence of healthcare-associated infections in acute care hospitals in Jacksonville, Florida. *Infect Control Hosp Epidemiol* 2012;33:283–291.
  18. European Centre for Disease Prevention and Control (ECDC). Point prevalence survey of healthcare-associated infections and antimicrobial use in European acute care hospitals. Stockholm: ECDC, 2013.
  19. Pellizzer G, Mantoan P, Timillero L, et al. Prevalence and risk factors for nosocomial infections in hospitals of the Veneto region, north-eastern Italy. *Infection* 2008;36:112–119.
  20. Durando P, Icardi G, Ansaldi F, et al. Surveillance of hospital-acquired infections in Liguria, Italy: results from a regional prevalence study in adult and paediatric acute-care hospitals. *J Hosp Infect* 2009;71:81–87.
  21. Pittet D, Harbarth S, Ruef C, et al. Prevalence and risk factors for nosocomial infections in four university hospitals in Switzerland. *Infect Control Hosp Epidemiol* 1999;20:37–42.
  22. Sax H, Hugonnet S, Harbarth S, Herrault P, Pittet D. Variation in nosocomial infection prevalence according to patient care setting: a hospital-wide survey. *J Hosp Infect* 2001;48:27–32.
  23. Sax H, Pittet D. Interhospital differences in nosocomial infection rates: importance of case-mix adjustment. *Arch Intern Med* 2002;162:2437–2442.
  24. Sax H. Nationwide surveillance of nosocomial infections in Switzerland—methods and results of the Swiss Nosocomial Infection Prevalence Studies (SNIP) in 1999 and 2002 [in German]. *Ther Umsch* 2004;61:197–203.
  25. Muhlemann K, Franzini C, Aepli C, et al. Prevalence of nosocomial infections in Swiss children's hospitals. *Infect Control Hosp Epidemiol* 2004;25:765–771.
  26. Weinstein JW, Mazon D, Pantelick E, Reagan-Cirincione P, Dembry LM, Hierholzer WJ Jr. A decade of prevalence surveys in a tertiary-care center: trends in nosocomial infection rates, device utilization, and patient acuity. *Infect Control Hosp Epidemiol* 1999;20:543–548.
  27. Vaque J, Rossello J, Trilla A, et al. Nosocomial Infections Prevalence Study in Spain. Nosocomial infections in Spain: results of five nationwide serial prevalence surveys (EPINE Project, 1990 to 1994). *Infect Control Hosp Epidemiol* 1996;17:293–297.
  28. Frankart L, Copin P, Alexiou A, Henry N, Sauvan V, Pittet D. Prevalence of nosocomial infections in a university hospital: distribution, predisposing factors and diagnostic indices [in French]. *Schweiz Med Wochenschr* 1998;128:1973–1983.
  29. McCabe WR, Jackson GG. Gram-negative bacteremia. I. Etiology and ecology. *Arch Intern Med* 1962;110:847–853.
  30. Charlson ME, Pompei P, Ales KL, MacKenzie CR. A new method of classifying prognostic comorbidity in longitudinal studies: development and validation. *J Chron Dis* 1987;40:373–383.
  31. Freeman J, Hutchison GB. Prevalence, incidence and duration. *Am J Epidemiol* 1980;112:707–723.
  32. Rhame FS, Sudderth WD. Incidence and prevalence as used in the analysis of the occurrence of nosocomial infections. *Am J Epidemiol* 1981;113:1–11.
  33. Gastmeier P, Brauer H, Sohr D, et al. Converting incidence and prevalence data of nosocomial infections: results from eight hospitals. *Infect Control Hosp Epidemiol* 2001;22:31–34.
  34. Eriksen HM, Iversen BG, Aavitsland P. Prevalence of nosocomial infections and use of antibiotics in long-term care facilities in Norway, 2002 and 2003. *J Hosp Infect* 2004;57:316–320.
  35. Eikelenboom-Boskamp A, Cox-Claessens JH, Boom-Poels PG, Drabbe MI, Koopmans RT, Voss A. Three-year prevalence of healthcare-associated infections in Dutch nursing homes. *J Hosp Infect* 2011;78:59–62.
  36. Rosenthal VD, Bijie H, Maki DG, et al. International Nosocomial Infection Control Consortium (INICC) report, data summary of 36 countries, for 2004–2009. *Am J Infect Control* 2012;40:396–407.
  37. Dudeck MA, Horan TC, Peterson KD, et al. National Healthcare Safety Network (NHSN) report, data summary for 2010, device-associated module. *Am J Infect Control* 2011;39:798–816.
  38. Gastmeier P, Sohr D, Schwab F, et al. Ten years of KISS: the most important requirements for success. *J Hosp Infect* 2008;70(suppl 1):11–16.
  39. Zingg W, Sax H, Inan C, et al. Hospital-wide surveillance of catheter-related bloodstream infection: from the expected to the unexpected. *J Hosp Infect* 2009;73:41–46.
  40. Hopmans TE, Blok HE, Troelstra A, Bonten MJ. Prevalence of hospital-acquired infections during successive surveillance surveys conducted at a university hospital in the Netherlands. *Infect Control Hosp Epidemiol* 2007;28:459–465.
  41. Sartor C, Sambuc R, Bimar MC, Gulian C, De Micco P. Prevalence surveys of nosocomial infections using a random sampling method in Marseille hospitals. *J Hosp Infect* 1995;29:209–216.
  42. Gastmeier P, Sohr D, Rath A, et al. Repeated prevalence investigations on nosocomial infections for continuous surveillance. *J Hosp Infect* 2000;45:47–53.
  43. Ebnother C, Tanner B, Schmid F, La Rocca V, Heinzer I, Brengener T. Impact of an infection control program on the prevalence of nosocomial infections at a tertiary care center in Switzerland. *Infect Control Hosp Epidemiol* 2008;29:38–43.
  44. Kallel H, Bahoul M, Ksibi H, et al. Prevalence of hospital-acquired infection in a Tunisian hospital. *J Hosp Infect* 2005;59:343–347.
  45. Atif ML, Bezzaoucha A, Mesbah S, Djellato S, Boubechou N, Bellouni R. Evolution of nosocomial infection prevalence in

- an Algeria university hospital (2001 to 2005) [in French]. *Med Mal Infect* 2006;36:423–428.
46. Jroundi I, Khoudri I, Azzouzi A, et al. Prevalence of hospital-acquired infection in a Moroccan university hospital. *Am J Infect Control* 2007;35:412–416.
  47. Razine R, Azzouzi A, Barkat A, et al. Prevalence of hospital-acquired infections in the university medical center of Rabat, Morocco. *Int Arch Med* 2012;5:26.
  48. El Rhazi K, Elfakir S, Berraho M, et al. Prevalence and risk factors for nosocomial infections in Hassan II University Hospital, Fes, Morocco [in French]. *East Mediterr Health J* 2007;13:56–63.
  49. Ogwang M, Paramatti D, Molteni T, et al. Prevalence of hospital-associated infections can be decreased effectively in developing countries. *J Hosp Infect* 2013;84:138–142.
  50. Jepsen OB, Jensen LP, Zimakoff J, et al. Prevalence of infections and use of antibiotics among hospitalized patients in Mauritius: a nationwide survey for the planning of a national infection control programme. *J Hosp Infect* 1993;25:271–278.
  51. Danchaivijitr S, Judaeng T, Sripalakij S, Naksawas K, Plipat T. Prevalence of nosocomial infection in Thailand 2006. *J Med Assoc Thai* 2007;90:1524–1529.
  52. Hughes AJ, Ariffin N, Huat TL, et al. Prevalence of nosocomial infection and antibiotic use at a university medical center in Malaysia. *Infect Control Hosp Epidemiol* 2005;26:100–104.
  53. Danchaivijitr S, Chokloikaew S. A national prevalence study on nosocomial infections 1988. *J Med Assoc Thai* 1989;72(suppl 2):1–6.
  54. Danchaivijitr S, Tangtrakool T, Chokloikaew S. The Second Thai National Prevalence Study on Nosocomial Infections 1992. *J Med Assoc Thai* 1995;78(suppl 2):S67–S72.
  55. Danchaivijitr S, Tangtrakool T, Waitayapichet S, Chokloikaew S. Efficacy of hospital infection control in Thailand 1988–1992. *J Hosp Infect* 1996;32:147–153.
  56. Thu TA, Hung NV, Quang NN, et al. A point-prevalence study on healthcare-associated infections in Vietnam: public health implications. *Infect Control Hosp Epidemiol* 2011;32:1039–1041.
  57. Stoesser N, Emary K, Soklin S, et al. The value of intermittent point-prevalence surveys of healthcare-associated infections for evaluating infection control interventions at Angkor Hospital for Children, Siem Reap, Cambodia. *Trans R Soc Trop Med Hyg* 2013;107:248–253.
  58. Ider BE, Clements A, Adams J, Whitby M, Muugolog T. Prevalence of hospital-acquired infections and antibiotic use in two tertiary Mongolian hospitals. *J Hosp Infect* 2010;75:214–219.
  59. Askarian M, Yadollahi M, Assadian O. Point prevalence and risk factors of hospital acquired infections in a cluster of university-affiliated hospitals in Shiraz, Iran. *J Infect Public Health* 2012;5:169–176.
  60. Lahsaeizadeh S, Jafari H, Askarian M. Healthcare-associated infection in Shiraz, Iran 2004–2005. *J Hosp Infect* 2008;69:283–287.
  61. Xie DS, Xiong W, Xiang LL, et al. Point prevalence surveys of healthcare-associated infection in 13 hospitals in Hubei Province, China, 2007–2008. *J Hosp Infect* 2010;76:150–155.
  62. Azzam R, Dramaix M. A one-day prevalence survey of hospital-acquired infections in Lebanon. *J Hosp Infect* 2001;49:74–78.
  63. Duerink DO, Roeshadi D, Wahjono H, et al. Surveillance of healthcare-associated infections in Indonesian hospitals. *J Hosp Infect* 2006;62:219–229.
  64. Abussaud MJ. Prevalence of nosocomial infections in a Saudi Arabian teaching hospital. *J Hosp Infect* 1991;17:235–238.
  65. McLaws ML, Gold J, King K, Irwig LM, Berry G. The prevalence of nosocomial and community-acquired infections in Australian hospitals. *Med J Aust* 1988;149:582–590.
  66. Graves N, Nicholls TM, Wong CG, Morris AJ. The prevalence and estimates of the cumulative incidence of hospital-acquired infections among patients admitted to Auckland District Health Board Hospitals in New Zealand. *Infect Control Hosp Epidemiol* 2003;24:56–61.
  67. Nicholls TM, Morris AJ. Nosocomial infection in Auckland Healthcare hospitals. *N Z Med J* 1997;110:314–316.
  68. Moro ML, Stazi MA, Marasca G, Greco D, Zampieri A. National prevalence survey of hospital-acquired infections in Italy, 1983. *J Hosp Infect* 1986;8:72–85.
  69. Bernander S, Hambræus A, Myrback KE, Nystrom B, Sundelof B. Prevalence of hospital-associated infections in five Swedish hospitals in November 1975. *Scand J Infect Dis* 1978;10:66–70.
  70. Jepsen OB, Mortensen N. Prevalence of nosocomial infection and infection control in Denmark. *J Hosp Infect* 1980;1:237–244.
  71. Lanini S, Jarvis WR, Nicastrì E, et al. Healthcare-associated infection in Italy: annual point-prevalence surveys, 2002–2004. *Infect Control Hosp Epidemiol* 2009;30:659–665.
  72. Signorelli C, D'Alessandro D, Collina D, Fara GM. Prevalence survey of nosocomial infections in a paediatric hospital. *J Hosp Infect* 1991;18:139–143.
  73. Pavia M, Bianco A, Viggiani NM, Angelillo IF. Prevalence of hospital-acquired infections in Italy. *J Hosp Infect* 2000;44:135–139.
  74. Di Pietrantonj C, Ferrara L, Lomolino G. Multicenter study of the prevalence of nosocomial infections in Italian hospitals. *Infect Control Hosp Epidemiol* 2004;25:85–87.
  75. Meers PD, Ayliffe GAJ, Emmerson AM, et al. Report on the National Survey of Infection in Hospitals, 1980. *J Hosp Infect* 1981;2:1–53.
  76. Mertens R, Kegels G, Stroobant A, et al. The national prevalence survey of nosocomial infections in Belgium, 1984. *J Hosp Infect* 1987;9:219–229.
  77. Gordts B, Vrijens F, Hulstaert F, Devriese S, Van de Sande S. The 2007 Belgian national prevalence survey for hospital-acquired infections. *J Hosp Infect* 2010;75:163–167.
  78. Aavitsland P, Stormark M, Lystad A. Hospital-acquired infections in Norway: a national prevalence survey in 1991. *Scand J Infect Dis* 1992;24:477–483.
  79. Scheel O, Stormark M. National prevalence survey on hospital infections in Norway. *J Hosp Infect* 1999;41:331–335.
  80. Lyytikäinen O, Kanerva M, Agthe N, Mottonen T, Ruutu P. Healthcare-associated infections in Finnish acute care hospitals: a national prevalence survey, 2005. *J Hosp Infect* 2008;69:288–294.
  81. Astagneau P, Fleury L, Leroy S, et al. Cost of antimicrobial treatment for nosocomial infections based on a French prevalence survey. *J Hosp Infect* 1999;42:303–312.
  82. The French Prevalence Survey Study Group. Prevalence of nosocomial infections in France: results of the nationwide survey in 1996. *J Hosp Infect* 2000;46:186–193.
  83. Floret N, Bailly P, Bertrand X, et al. Results from a four-year study on the prevalence of nosocomial infections in Franche-

- Comte: attempt to rank the risk of nosocomial infection. *J Hosp Infect* 2006;63:393–398.
84. Lietard C, Lejeune B, Metzger MH, Thiolet JM, Coignard B. National point prevalence survey of healthcare-associated infections: results for people aged 65 and older, France, 2006. *J Am Geriatr Soc* 2011;59:763–765.
  85. Vincent JL, Bihari DJ, Suter PM, et al; EPIC International Advisory Committee. The prevalence of nosocomial infection in intensive care units in Europe: results of the European Prevalence of Infection in Intensive Care (EPIC) Study. *JAMA* 1995; 274:639–644.
  86. Humphreys H, Newcombe RG, Enstone J, et al. Four country healthcare associated infection prevalence survey 2006: risk factor analysis. *J Hosp Infect* 2008;69:249–257.
  87. Fitzpatrick F, McIlvenny G, Oza A, et al. Hospital infection society prevalence survey of Healthcare Associated Infection 2006: comparison of results between Northern Ireland and the Republic of Ireland. *J Hosp Infect* 2008;69:265–273.
  88. Emmerson AM, Enstone JE, Griffin M, Kelsey MC, Smyth ET. The Second National Prevalence Survey of infection in hospitals: overview of the results. *J Hosp Infect* 1996;32:175–190.
  89. Smyth ET, McIlvenny G, Enstone JE, et al. Four country healthcare associated infection prevalence survey 2006: overview of the results. *J Hosp Infect* 2008;69:230–248.
  90. Reilly J, Cairns S, Fleming S, et al. Results from the second Scottish national prevalence survey: the changing epidemiology of healthcare-associated infection in Scotland. *J Hosp Infect* 2012;82:170–174.
  91. Reilly JSS, Allardice G, Noone A, Robertson C, Walker A, Coubrough S. *NHS Scotland National HAI Prevalence Survey: Health Protection Scotland: Final Report*. Glasgow, 2007.
  92. Vaque J, Rossello J, Arribas JL; EPINE Working Group. Prevalence of nosocomial infections in Spain: EPINE study 1990–1997. *J Hosp Infect* 1999;43(suppl):S105–S111.
  93. Ruden H, Gastmeier P, Daschner FD, Schumacher M. Nosocomial and community-acquired infections in Germany: summary of the results of the First National Prevalence Study (NIDEP). *Infection* 1997;25:199–202.
  94. Gastmeier P, Kampf G, Wischniewski N, et al. Prevalence of nosocomial infections in representative German hospitals. *J Hosp Infect* 1998;38:37–49.
  95. Kampf G, Wischniewski N, Schulgen G, Schumacher M, Daschner F. Prevalence and risk factors for nosocomial lower respiratory tract infections in German hospitals. *J Clin Epidemiol* 1998;51:495–502.
  96. van der Kooi TI, Mannien J, Wille JC, van Benthem BH. Prevalence of nosocomial infections in the Netherlands, 2007–2008: results of the first four national studies. *J Hosp Infect* 2010;75: 168–172.
  97. Valinteliene R, Jurkuvenas V, Jepsen OB. Prevalence of hospital-acquired infection in a Lithuanian hospital. *J Hosp Infect* 1996; 34:321–329.
  98. Dumpis U, Balode A, Vigante D, et al. Prevalence of nosocomial infections in two Latvian hospitals. *Euro Surveill* 2003; 8:73–78.
  99. Raka L, Zoutman D, Mulliqi G, et al. Prevalence of nosocomial infections in high-risk units in the university clinical center of Kosova. *Infect Control Hosp Epidemiol* 2006;27:421–423.
  100. Markovic-Denic L, Jankovic S, Bojanic J, Maksimovic N. The prevalence study of hospital-acquired infections at different surgical departments in Banjaluka [in Serbian]. *Srp Arh Celok Lek* 2006;134:229–233.
  101. Faria S, Sodano L, Gjata A, et al. The first prevalence survey of nosocomial infections in the University Hospital Centre ‘Mother Teresa’ of Tirana, Albania. *J Hosp Infect* 2007;65:244–250.
  102. Gikas A, Padiaditis J, Papadakis JA, et al. Prevalence study of hospital-acquired infections in 14 Greek hospitals: planning from the local to the national surveillance level. *J Hosp Infect* 2002;50:269–275.
  103. Esen S, Leblebicioglu H. Prevalence of nosocomial infections at intensive care units in Turkey: a multicentre 1-day point prevalence study. *Scand J Infect Dis* 2004;36:144–148.
  104. Metintas S, Akgun Y, Durmaz G, Kalyoncu C. Prevalence and characteristics of nosocomial infections in a Turkish university hospital. *Am J Infect Control* 2004;32:409–413.
  105. Sramova H, Bartonova A, Bolek S, Krecmerova M, Subertova V. National prevalence survey of hospital-acquired infections in Czechoslovakia. *J Hosp Infect* 1988;11:328–334.
  106. Klavs I, Bufon Luznik T, Skerl M, et al. Prevalence of and risk factors for hospital-acquired infections in Slovenia: results of the first national survey, 2001. *J Hosp Infect* 2003;54:149–157.
  107. Campins M, Vaque J, Rossello J, et al. Nosocomial infections in pediatric patients: a prevalence study in Spanish hospitals. EPINE Working Group. *Am J Infect Control* 1993;21:58–63.
  108. Nicastrì E, Petrosillo N, Martini L, Larosa M, Gesu GP, Ippolito G. Prevalence of nosocomial infections in 15 Italian hospitals: first point prevalence study for the INF-NOS project. *Infection* 2003;31(suppl 2):10–15.
  109. Lizioli A, Privitera G, Alliata E, et al. Prevalence of nosocomial infections in Italy: result from the Lombardy survey in 2000. *J Hosp Infect* 2003;54:141–148.
  110. Zotti CM, Messori Ioli G, Charrier L, et al. Hospital-acquired infections in Italy: a region wide prevalence study. *J Hosp Infect* 2004;56:142–149.
  111. Stevens GP, Jacobson JA, Burke JP. Changing patterns of hospital infections and antibiotic use: prevalence surveys in a community hospital. *Arch Intern Med* 1981;141:587–592.
  112. Avila-Figueroa C, Cashat-Cruz M, Aranda-Patron E, et al. Prevalence of nosocomial infections in children: survey of 21 hospitals in Mexico [in Spanish]. *Salud Publica Mex* 1999;41(suppl 1):S18–S25.
  113. Gravel D, Taylor G, Ofner M, et al. Point prevalence survey for healthcare-associated infections within Canadian adult acute-care hospitals. *J Hosp Infect* 2007;66:243–248.
  114. Rezende EM, Couto BR, Starling CE, Modena CM. Prevalence of nosocomial infections in general hospitals in Belo Horizonte. *Infect Control Hosp Epidemiol* 1998;19:872–876.
  115. Guanche Garcell H, Nunez Labrador L, Baxter Campana M, et al. Prevalence of nosocomial infections in University Hospital of Habana [in Spanish]. *An Med Interna* 2006;23:269–271.
  116. Izquierdo-Cubas F, Zambrano A, Frometa I, et al. National prevalence of nosocomial infections: Cuba 2004. *J Hosp Infect* 2008;68:234–240.
  117. Durlach R, McIlvenny G, Newcombe RG, et al. Prevalence survey of healthcare-associated infections in Argentina; comparison with England, Wales, Northern Ireland and South Africa. *J Hosp Infect* 2012;80:217–223.
  118. Suetens C, Ammon A, Weist K, Sodano L, Monnet DL. Review of methods of national prevalence surveys of healthcare as-



- sociated infections in 17 European countries. In: *European Congress of Clinical Microbiology and Infectious Diseases (ECCMID)*. May 16–19, 2009; Helsinki. Abstract P624.
119. Haley RW, Quade D, Freeman HE, Bennett JV. The SENIC Project: study on the efficacy of nosocomial infection control (SENIC Project); summary of study design. *Am J Epidemiol* 1980;111:472–485.
  120. World Health Organization (WHO). *Surveillance, Control and Prevention of Hospital-Acquired (Nosocomial) Infections: Report of an Advisory Group*. Geneva: WHO, 1981.
  121. Nosocomial Infection Control Group of Thailand. Definitions of nosocomial infections. *J Med Assoc Thai* 1988;71:58–63.
  122. EPINE Working Group. Prevalence of hospital-acquired infections in Spain. *J Hosp Infect* 1992;20:1–13.
  123. Department of Health and Human Services. *Outline for Surveillance and Control of Nosocomial Infections*. Appendix 11. Atlanta: Centers for Disease Control, US Public Health Service, 1988.
  124. Emori TG, Culver DH, Horan TC, et al. National Nosocomial Infections Surveillance System (NNIS): description of surveillance methods. *Am J Infect Control* 1991;19:19–35.
  125. Quenon JL, Gottot S, Duneton P, et al. Enquête nationale de prévalence des infections nosocomiales en France. *Bull Epidemiol Hebdo* 1993;39:179–180.